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	- 1		EPO; JPO;	
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2	3	(Sheppard NEAR Michael) and endostatin	USPAT;	2004/05/24 15:07
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			EPO; JPO;	
			DERWENT	
4	11	Tong NEAR Xiao	USPAT;	2004/05/24 15:08
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5	1712	endostatin	USPAT;	2004/05/24 15:08
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7	7	endostatin SAME (cani\$3 or dog)	USPAT;	2004/05/24 15:10
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8	/	(US-20030139365-\$ or US-20030158099-\$ or US-20030059417-\$).did. or (EP-1197550-\$ or	EPO; JPO	2004/05/24 15.11
		EP-1191036-\$).did. or (JP-2003000268-\$ or	EFO, UIO	
		JP-2002355056-\$).did.		
	- 2	f = = = : :	USPAT;	2003/09/15 14:47
_		20030133303	US-PGPUB;	2003,03,13 14.47
			EPO; JPO;	
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	55	Gillies NEAR Stephen	USPAT;	2003/09/15 14:54
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(FILE 'HOME' ENTERED AT 15:11:43 ON 24 MAY 2004)

FILE 'MEDLINE, CAPLUS' ENTERED AT 15:12:01 ON 24 MAY 2004

L1 3 S ENDOSTATIN (L) (CANI? OR DOG)

L2 3 DUP REM L1 (0 DUPLICATES REMOVED)

E SPEPPARD MICHAEL?/AU

E SHEPPARD MICHAEL?/AU

L3 9 S E1

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L2 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2002:237903 CAPLUS

DN 136:259025

TI cDNA encoding **endostatin** and its use as inhibitor of cancer and angiogenesis-related disorders in **dogs** 

SO Eur. Pat. Appl., 56 pp.

CODEN: EPXXDW

IN Sheppard, Michael G.; Tong, Xiao AB The present invention relates to

The present invention relates to canine endostatin genes and polypeptides associated as inhibitors of angiogenesis-related disorders, such as cancer. The invention encompasses endostatin nucleic acids, recombinant DNA mols., cloned genes or degenerate variants thereof, endostatin gene products and antibodies directed against such gene products, cloning vectors containing mammalian endostatin gene mols., and hosts that have been genetically engineered to express such mols. In a further embodiment the said endostatin is not from chicken, human or mouse. The invention further relates to methods for the identification of compds. that modulate the expression of endostatin genes and gene products and to using such compds. as therapeutic agents in the treatment of angiogenesis-related disorders, e.g., cancer. The invention also relates to methods for the diagnostic evaluation, genetic testing and prognosis of angiogenesis-related disorders, e.g., cancer, and to methods and compns. for the treatment these disorders. Angiogenesis-related disorders include angiogenesis-dependent cancers comprising solid tumors and blood borne tumors such as leukemias, tumor metastases, benign tumors comprising hemaniomas, acoustic neuromas, neurofibromas, trachomas, and pyogenic granulomas. Also included are rheumatoid arthritis, psoriasis, ocular angiogenic diseases comprising diabetic retinopathy, retinopathy of prematurity, macular degeneration, corneal graft rejection, neo-vascular qlaucoma, retrolental fibroplasia, and rubeosis. Osler-Webber syndrome, myocardial angiogenesis, plaque neovascularization, telangiectasia, hemophiliac joints, angiofibroma, wound granulation, coronary collaterals, cerebral collaterals, and arteriovenous malformations are also included. Other diseases include ischemic limb angiogenesis, diabetic neovascularization, macular degeneration, fractures, vasculogenesis, hematopoiesis, ovulation, menstruation and placentation.

	PATENT NO.	KIND DATE	APPLICATION NO.	DATE
PI	EP 1191036	A2 20020327	EP 2001-307224	20010824
	EP 1191036	A3 20020703		
	R: AT, BE,	CH, DE, DK, ES,	FR, GB, GR, IT, LI, LU,	NL, SE, MC, PT,
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	JP 2003000268	A2 20030107	JP 2001-254697	20010824
	US 2003158099	A1 20030821	US 2001-938391	20010824

- L2 ANSWER 2 OF 3 MEDLINE on STN
- AN 2002487981 MEDLINE
- TI Endostatin concentrations in healthy dogs and dogs with selected neoplasms.
- SO Journal of veterinary internal medicine / American College of Veterinary Internal Medicine, (2002 Sep-Oct) 16 (5) 565-9.

  Journal code: 8708660. ISSN: 0891-6640.
- AU Rossmeisl John H Jr; Bright Patricia; Tamarkin Lawrence; Simpson Byron W; Troy Gregory C; Hueston William; Ward Daniel L
- AB Endostatin prevents angiogenesis and tumor growth by inhibiting

endothelial cell proliferation and migration. The purpose of this study was to determine serum endostatin concentrations in 53 healthy dogs and in 38 dogs with confirmed malignant neoplasms. Endostatin concentration was determined with a competitive enzymatic immunoassay (EIA) with rabbit polyclonal antibody generated against a recombinant canine endostatin protein. Both the presence of cancer and increasing age were associated with increased serum concentration of endostatin. Endostatin concentration in healthy dogs was 87.7 +/- 3.5 ng/mL. Upper and lower limits of the reference range for serum endostatin concentration in healthy dogs were 60 and 113 ng/mL. Dogs with lymphoma (LSA) and hemangiosarcoma (HSA) had endostatin concentrations of 107 +/- 9.3 ng/mL. In conclusion, this study demonstrates that endostatin can be quantified in dogs and that endostatin concentrations are high in dogs with HSA and LSA.

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L2 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN
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AN 2000:144909 CAPLUS

DN 132:190503

TI Expression and export of angiostatin and endostatin as immunofusins

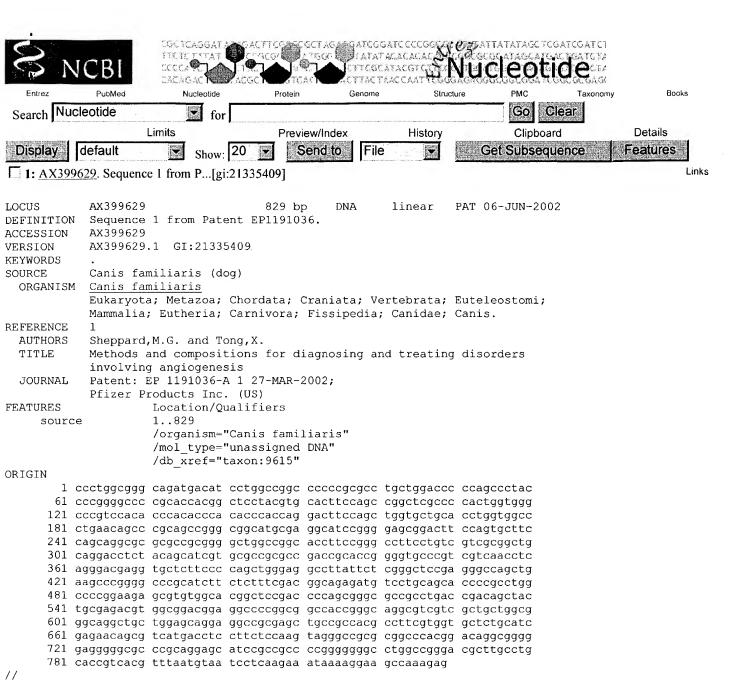
SO PCT Int. Appl., 68 pp.

CODEN: PIXXD2

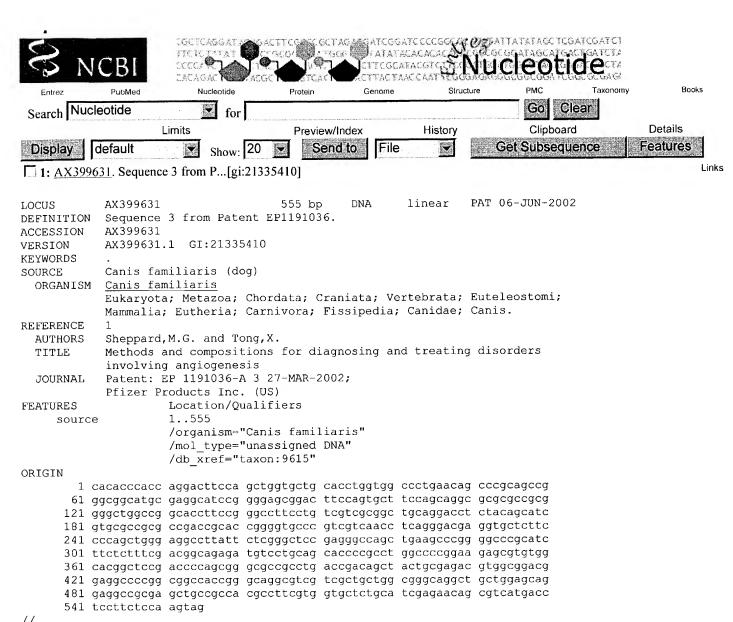
IN Lo, Kin-Ming; Li, Yue; Gillies, Stephen D.

AB Disclosed are nucleotide sequences, for example, DNA or RNA sequences, which encode an Ig Fc-angiogenesis inhibitor fusion protein. The angiogenesis inhibitors can be angiostatin, endostatin, a plasminogen fragment having angiostatin activity, or a collagen XVIII fragment having endostatin activity. The nucleotide sequences can be inserted into a suitable expression vector and expressed in mammalian cells. Also disclosed is a family of Ig Fc-angiogenesis inhibitor fusion proteins that can be produced by expression of such nucleotide sequences. Also disclosed are methods using such nucleotide sequences and fusion proteins for treating conditions mediated by angiogenesis. When C57/BL6 mice with implanted Lewis lung tumors are injected with 720 μg human Fc-human angiostatin fusion protein per mouse, the protein had a circulating half-life of about 32 h, and Western anal. shows that >90% of the fusion protein remains as an intact mol. in circulation.

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PATENT NO.
                       KIND DATE
                                               APPLICATION NO.
PΙ
     WO 2000011033
                         A2
                               20000302
                                               WO 1999-US19329 19990825
     WO 2000011033
                         A3
                               20000622
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